

## TITLE: BONE STIMULATING FACTOR

[0001] This application is a continuation of prior Serial No. 09/229,304 filed January 13, 1999, now United States Patent No. \_\_\_\_\_ issued, which application is a continuation-in-part of prior application Serial No. 09/048,058 filed March 26, 1998, which application is a continuation application of prior application Serial No. PCT/CA 96/00653 filed September 26, 1996, which application claims priority from United States Provisional Patent Application Serial No. 60/004,314 filed September 16, 1995, all of which applications are incorporated herein by reference.

[0002] The present invention relates to polypeptides which stimulate bone growth.

[0003] Understanding of issues related to bone growth and strength has progressed over the years, a summary being provided in, for example, international patent application No. PCT/CA 94/00144, published on September 15, 1994 under WO 94/20615, United States Patent No. 5,320,970 and European patent application No. 92 302 446, published under 505 210 on September 23, 1992, the contents of which applications are incorporated herein by reference.

[0004] By way of background to the present invention, described below, neutrophil-activating peptide (NAP-2; SEQ ID NO:1) and a variant of NAP-2, here termed "NAP-2V" (SEQ ID NO:2) have been known for some time (Walz, A., and M. Baggiolini, (1989) *Biochem. Biophys. Res. Commun.* 159:969). British Patent No. 2 231 872 (British Patent No. 2 231 872. Inventors: M. Baggiolini, K.J. Clemetson, and A. Walz. Published June 14, 1990.), describes the amino acid sequence of NAP-2 and three apparently naturally occurring variants, including NAP-2V. The other two variants have an additional four (SEQ ID NO:3) and three (SEQ ID NO:4) amino acids at the N-terminal of the NAP-2 sequence. NAP-2 is a subsequence of  $\beta$ -thromboglobulin ( $\beta$ -TG; SEQ ID NO:5) which has an additional eleven amino acids at the N-terminal end.  $\beta$ -TG is itself a subsequence of conn ective

tissue-activating peptide (CTAP-III; SEQ ID NO:6) which has an additional four amino acids at the N-terminal. CTAP-III is a subsequence of platelet basic protein (PBP; SEQ ID NO:7) which has an additional nine amino acids at the N-terminal.

[0005] NAP-2 along with interleukin-8 (human IL-8; SEQ ID NO:8; porcine IL-8 SEQ ID NO:9) and melanoma growth-stimulating activity (MGSA) have been assigned to a subfamily known as the  $\alpha$ -chemokines. The  $\alpha$ -chemokines have in common with each other four cysteine residues at highly conserved positions, which enclose the core region of the molecules, as described by Brandt *et al* (Ehlert, J.E., F. Peterson, M.H.G. Kubbuta, J. Gerdes, H.-D. Flad, and E. Brandt, (1995) *J. Biol. Chem.* 270:6338). Brandt *et al.* found an apparently naturally occurring C-terminus truncated variant of NAP-2, lacking the last four amino acids of NAP-2, that displays enhanced increase in potency to stimulate neutrophil degranulation. Brandt *et al.* also synthesized variants lacking the final one, two, three, five and six amino acids of the C-terminus of NAP-2. All of these C-truncated polypeptides exhibited a moderate increase in potency over NAP-2 with the exception of the sequence having only the first sixty-four amino acids of NAP-2. Brandt *et al.* discussed the possible significance of the sequence modifications with respect to the structure of NAP-2 and its function.

[0006] Platelet factor 4 (PF4; SEQ ID NO:10) is a seventy amino acid polypeptide (Hermanson, M., G. Schmer and K. Kurachi, (1977) *J. Biol. Chem.* 252:6276; Morgan, F.J., G.S. Begg, C.N. Chesterman, (1979) *Thromb. Haemost.* 42:1652). PF4 has been shown to inhibit proliferation of two osteoblastic osteosarcoma cell lines, Saos-2 and G-292 (United States Patent No. 5,304,542. Inventor: D.M. Tatakis. Issued April 19, 1994). Indomethacin apparently did not affect PF4-induced inhibition of the cell proliferation. Particular fragments, PF4(58-70), PF4(47-70) and monomeric low-affinity PF4 (LAPF4), which is 50% homologous to PF4 and contains an  $\alpha$ -helical C-terminus were also suggested as being useful. PF4 and such related polypeptides were thus described as being useful in a method for

inhibiting proliferation of osteoblasts, in among other things, humans suffering from osteoporosis.

[0007] The first 70 amino acids of NAP-2V and the sequence of PF4 are about 51% homologous and the positions of the four cysteine residues are conserved between the two polypeptides.

[0008] It has now been shown that NAP-2, NAP-2V, as well as subsequences of NAP-2V also show bone stimulatory effects, while certain subsequences do not display bone stimulatory activity. NAP-2V-(1-26) (SEQ ID NO:11) and NAP-2V-(13-26; gln<sup>25</sup>→glu<sup>25</sup>) (SEQ ID NO:12) were found to increase the observed bone apposition rate, the latter of the two being more potent than the former. NAP-2V-(10-26) (SEQ ID NO:13) appeared to cause a small increase in the observed bone apposition rate, although the statistical significance of the observed increase was questionable. Protected versions of NAP-2V-(13-26; gln<sup>25</sup>→glu<sup>25</sup>) (SEQ ID NO:17) and of NAP-2V-(15-22) (SEQ ID NO:18) were also found to have bone stimulatory activity. NAP-2V-(11-26) (SEQ ID NO:14) and NAP-2V-(12-26) (SEQ ID NO:15) were found to have no effect on observed bone mineral apposition rate.

[0009] The invention thus includes a compound which promotes bone growth in mammals, comprising an amino acid sequence which consists essentially of:

[0010] (i) up to 69 consecutive amino acids of the sequence corresponding to SEQ ID NO:2 with (a) from 6 to about 14 amino acids deleted from the N-terminus of SEQ ID NO:2, or (b) 7 to about 53 amino acids deleted from the C-terminus of SEQ ID NO:2, or both (a) and (b) and wherein the sequence includes no cysteine residues or at least two cysteine residues;

[0011] (ii) a variant of a polypeptide of (i) containing a plurality of said amino acid sequences;

[0012] (iii) a conservatively substituted variant of a polypeptide of (i) or (ii);  
or

[0013] (iv) a functionally equivalent homologue of (i), (ii) or (iii).

[0014] In another aspect, the invention includes a polypeptide in which the amino acid sequence consists essentially of an amino acid sequence corresponding to SEQ ID NO:11, or corresponding to SEQ ID NO:11 with from 6 to about 14 amino acids deleted from the N-terminus of SEQ ID NO:11, or corresponding to SEQ ID NO:11 with from 6 to about 12 amino acids deleted from the N-terminus of SEQ ID NO:11; or corresponding to SEQ ID NO:11 with from 6 to about 9 amino acids deleted from the N-terminus of SEQ ID NO:11; or a functionally equivalent homologue any of the foregoing polypeptides.

[0015] A polypeptide of the present invention can have an amino acid sequence which consists essentially of an amino acid sequence corresponding to SEQ ID NO:12, or SEQ ID NO:13, or SEQ ID NO:18, or a functionally equivalent homologue of any of these.

[0016] The present invention includes a compound "derived from" any of the preceding polypeptides.

[0017] The present invention thus includes a polypeptide wherein the amino acid sequence consists essentially of:

- [0018] (v) an amino acid sequence corresponding to SEQ ID NO:12;
- [0019] (vi) an amino acid sequence corresponding to SEQ ID NO:13;
- [0020] (vii) an amino acid sequence corresponding to SEQ ID NO:18;
- [0021] (viii) a variant of a polypeptide of (v), (vi) or (vii) containing a plurality of said amino acid sequences; or
- [0022] (ix) a conservatively substituted variant of a polypeptide of (v), (vi), (vii) or (viii).

[0023] In preferred embodiments, a polypeptide of the present invention that cysteine residues, includes two cysteine residues that are located at positions corresponding to the tenth and twelfth positions of SEQ ID NO:2.

[0024] A polypeptide of the present invention can have, if desired, or necessary, one or the other or both of the N-terminal amino acid and the C-terminal amino acid protected by a protecting group.

[0025] The polypeptide can be synthetic and the amino acid sequence can have a molecular weight in the range of from about 1000 to 4000 daltons; or

from about 1000 to about 3000; or from about 1000 to about 2000; or more preferably from about 1200 to about 1800.

[0026] In another aspect, the invention is a first polypeptide which promotes bone growth in mammals comprising a sequence of amino acids which consists essentially of up to 69 amino acids and is sufficiently duplicative of a second polypeptide such that the first polypeptide is encoded by a DNA that selectively hybridizes under stringent conditions with DNA encoding the second polypeptide. In this instance, the second polypeptide comprises an amino acid sequence which consists essentially of:

[0027] (i) up to 69 consecutive amino acids of the sequence corresponding to SEQ ID NO:2 with (a) from 6 to about 14 amino acids deleted from the N-terminus of SEQ ID NO:2, or (b) 7 to about 53 amino acids deleted from the C-terminus of SEQ ID NO:2, or both (a) and (b) and wherein the sequence includes no cysteine residues or at least two cysteine residues;

[0028] (ii) a variant of a polypeptide of (i) containing a plurality of said amino acid sequences;

[0029] (iii) a conservatively substituted variant of a polypeptide of (i) or (ii);  
or

[0030] (iv) a functionally equivalent homologue of (i), (ii) or (iii).

[0031] The DNA sequence of NAP-2V disclosed by Walz *et al.* (British Patent No. 2 231 872. Inventors: M. Baggiolini, K.J. Clemetson, and A. Walz. Published June 14, 1990. Neutrophil-activating peptide-2 and processes for the production of NAP-2, B-TG, CTAP-III and PBP) is identified here as SEQ ID NO:16.

[0032] The phrase "selectively hybridizes to" refers to a nucleic acid molecules that, under appropriately stringent hybridization conditions, hybridize, duplex or bind essentially only to each other when molecules having the predefined sequences (i.e., second polypeptide) are present in a preparation of DNA or RNA. For discussions of nucleic acid molecule design and annealing conditions, see, for example, Sambrook *et al.*, *Molecular Cloning: A Laboratory Manual* (2nd ed.), Vols. 1-3, Cold Spring Harbor Laboratory,

(1989) or Current Protocols in Molecular Biology, F. Ausubel et al., (ed.)

Greene Publishing and Wiley-Interscience, New York (1987).

[0033] It will, of course, be understood by those skilled in the art that portions of the nucleic acid sequence identified as SEQ ID NO:16 correspond to sequences coding for the polypeptides identified as SEQ ID NO:1, NO:11; NO:12 (Glu→Gln) or NO:13. For example, nucleic acids 37 through 78 of SEQ ID NO:16 encode the amino acid subsequence identified as SEQ ID NO:12 in which the penultimate amino acid of the subsequence is glutamine.

[0034] "Stringent hybridization conditions" takes on its common meaning to a person skilled in the art here. Appropriate stringency conditions which promote nucleic acid hybridization, for example, 6x sodium chloride/sodium citrate (SSC) at about 45°C are known to those skilled in the art. The following examples are found in Current Protocols in Molecular Biology, John Wiley & Sons, NY (1989), 6.3.1-6.3.6: For 50 ml of a first suitable hybridization solution, mix together 24 ml formamide, 12 ml 20x SSC, 0.5 ml 2 M Tris-HCl pH 7.6, 0.5 ml 100x Denhardt's solution, 2.5 ml deionized H<sub>2</sub>O, 10 ml 50% dextran sulfate, and 0.5 ml 10% SDS. A second suitable hybridization solution can be 1% crystalline BSA (fraction V), 1 mM EDTA, 0.5 M Na<sub>2</sub>HPO<sub>4</sub> pH 7.2, 7% SDS. The salt concentration in the wash step can be selected from a low stringency of about 2x SSC at 50 °C to a high stringency of about 0.2x SSC at 50°C. Both of these wash solutions may contain 0.1% SDS. In addition, the temperature in the wash step can be increased from low stringency conditions at room temperature, about 22°C to high stringency conditions, at about 65°C. The cited reference gives more detail, but appropriate wash stringency depends on degree of homology and length of probe. If homology is 100%, a high temperature (65°C to 75°C) may be used. If homology is low, lower wash temperatures must be used. However, if the probe is very short (<100bp), lower temperatures must be used even with 100% homology. In general, one starts washing at low temperatures (37°C to 40°C), and raises the temperature by 3-5°C intervals until background is low enough not to be a major factor in autoradiography.

**[0035]** Alternatively, such a first polypeptide is a sequence of amino acids sufficiently duplicative of a second polypeptide having an amino acid sequence corresponding to SEQ ID NO:11 up to 69 amino acids in length, or corresponding to SEQ ID NO:11 with from 6 to about 12 amino acids deleted from the N-terminus of SEQ ID NO:11, or corresponding to SEQ ID NO:11 with from 6 to about 9 amino acids deleted from the N-terminus of SEQ ID NO:11; wherein the sequence includes no cysteine residues or at least two cysteine residues; or a functionally equivalent homologue, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide. The second polypeptide can have, alternatively, or additionally 7 to about 53 amino acids deleted from the C-terminus of SEQ ID NO:2.

**[0036]** The first polypeptide can have a sequence of amino acids sufficiently duplicative of a second polypeptide having an amino acid sequence corresponding to SEQ ID NO:12 up to 69 amino acids in length; or a functionally equivalent homologue, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide.

**[0037]** The first polypeptide can have a sequence of amino acids sufficiently duplicative of a second polypeptide having an amino acid sequence corresponding to SEQ ID NO:11; or a conservatively substituted variant thereof, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide; or it can have a sequence of amino acids sufficiently duplicative of a second polypeptide comprising an amino acid sequence corresponding to SEQ ID NO:12; or a conservatively substituted variant thereof, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide; or it can have a sequence of amino acids sufficiently duplicative of a second polypeptide having an amino acid sequence corresponding to SEQ ID NO:13; or a conservatively substituted variant thereof, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second

polypeptide; or it can have a sequence of amino acids sufficiently duplicative of a second polypeptide having an amino acid sequence corresponding to SEQ ID NO:18; or a conservatively substituted variant thereof, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide.

[0038] A bone growth polypeptide of the invention can be of any suitable length, and particularly can be up to 8 amino acids in length, or up to 14 amino acids in length, or up to 20 amino acids in length, or up to 30 amino acids in length, or up to 40 amino acids in length, or up to 50 amino acids in length, or up to 60 amino acids or more in length.

[0039] The invention includes any number of chimeric bone stimulating factors made having an amino acid sequence of polypeptides of the present invention.

[0040] In another aspect, the invention is an agent for use in prevention and treatment of a bone reduction related disease which includes any polypeptide or polypeptides of the present invention as an active ingredient.

[0041] The invention is, alternatively, a pharmaceutical composition for promoting bone growth, having a therapeutically effective amount of a polypeptide or polypeptides of the present invention.

[0042] The invention includes a method of increasing bone growth in a mammal by administering a therapeutically effective amount of a polypeptide having an amino acid sequence of a polypeptide or polypeptides of the present invention.

[0043] The invention includes use of a polypeptide or polypeptides of the present invention for the treatment of osteoporosis. Alternatively, the use of a polypeptide or polypeptides can be to promote bone growth in a mammal.

[0044] The invention includes use of the polypeptide or polypeptides in the preparation of a medicament for use in promoting bone growth or the treatment of osteoporosis.

[0045] The invention includes a diagnostic kit for determining the presence of a polypeptide or polypeptides of the present invention. The kit can include an antibody to a said polypeptide(s) linked to a reporter system wherein the



reporter system produces a detectable response when a predetermined amount of the polypeptide(s) and the antibody are bound together.

[0046] The invention includes an antibody synthesized using a polypeptide consisting of an amino acid sequence identified as SEQ ID NO:11; SEQ ID NO:12; or SEQ ID NO:13 or a conservatively substituted variant thereof.

[0047] More generally, the invention includes an antibody which binds to a polypeptide or polypeptides of the present invention, synthesized using the polypeptide(s).

[0048] The invention includes an isolated DNA fragment which encodes the expression of any of the polypeptides of the present invention, and DNA which differs from the fragment due to the degeneracy of the genetic code.

[0049] The invention includes a vector comprising a DNA sequence which encodes the expression of any of any of the polypeptides of the present invention.

[0050] The invention includes a process for producing a polypeptide of the invention which includes the steps of:

[0051] a) preparing a DNA fragment containing a nucleotide sequence which encodes the polypeptide;

[0052] b) incorporating said DNA fragment into an expression vector to obtain a recombinant DNA fragment which contains the DNA fragment and is capable of undergoing replication;

[0053] c) transforming a host cell with the recombinant DNA fragment to isolate a transformant which can express the polypeptide; and

[0054] d) culturing the transformant to allow the transformant to produce the polypeptide and recovering the polypeptide from resulting cultured mixture.

[0055] In yet another aspect, the invention is a synthetic polypeptide up to 65 amino acids in length, having *in vivo* bone stimulatory activity in mammals, having an amino acid sequence which is at least about 11% conserved in relation to the amino acid sequence identified as SEQ ID NO:2 (i.e., contains 8 consecutive amino acids of the sequence identified as SEQ ID NO:2); a

conservatively substituted variant thereof; or a functionally equivalent homologue.

[0056] The synthetic polypeptide can also be at least about 11%, 12%, 13%, 15%, 16%, 17%, 19%, 22%, 25%, 28%, 31%, or 35% conserved in relation to the amino acid sequence identified as SEQ ID NO:2; a conservatively substituted variant thereof; or a functionally equivalent homologue.

[0057] Such a synthetic polypeptide can have at least 49 amino acids deleted from the sequence.

[0058] The polypeptide can have no cysteine residues or at least two cysteine residues.

[0059] The polypeptide can have a molecular weight in the range of from about 1000 to 4000.

[0060] In another aspect, the present invention is a first polypeptide having a sequence of amino acids sufficiently duplicative of a second polypeptide which includes an amino acid sequence of any of the synthetic polypeptides, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide.

[0061] The invention includes a chimeric bone stimulating factor comprising an amino acid sequence of any of the synthetic polypeptides.

[0062] An agent of the present invention for use in prevention and treatment of a bone reduction related disease can include one or more of the synthetic polypeptides.

[0063] As well, a pharmaceutical composition of the present invention for promoting bone growth, can include a therapeutically effective amount of a one or more of the synthetic polypeptides.

[0064] The invention includes a method of increasing bone growth in a mammal by administering a therapeutically effective amount of one or more of the synthetic polypeptides.

[0065] Such a synthetic polypeptide can be used for the treatment of osteoporosis or to promote bone growth in a mammal.

[0066] Such a synthetic polypeptide can be used in the preparation of a medicament for use in promoting bone growth or the treatment of osteoporosis.

[0067] The invention includes a diagnostic kit for determining the presence of one or more the synthetic polypeptides, the kit including antibody(ies) to a the polypeptide(s) linked to a reporter system wherein the reporter system produces a detectable response when a predetermined amount of the polypeptide(s) and the antibody(ies) are bound together.

[0068] The invention includes an antibody which binds to one or more of the synthetic polypeptides, synthesized using the polypeptide.

[0069] Further still, the invention includes an isolated DNA fragment which encodes the expression of any of the synthetic polypeptides, and DNA which differs from the fragment due to the degeneracy of the genetic code. The invention includes a vector which includes such a DNA sequence.

[0070] The invention includes a process for producing any of the synthetic polypeptides, which includes:

[0071] a) preparing a DNA fragment containing a nucleotide sequence which encodes a polypeptide;

[0072] b) incorporating said DNA fragment into an expression vector to obtain a recombinant DNA fragment which contains the DNA fragment and is capable of undergoing replication;

[0073] c) transforming a host cell with the recombinant DNA fragment to isolate a transformant which can express the polypeptide; and

[0074] d) culturing the transformant to allow the transformant to produce the polypeptide and recovering the polypeptide from resulting cultured mixture.

[0075] The invention includes a polypeptide exhibiting bone stimulatory activity in mammals, the polypeptide being up to 65 amino acids in length and having the sequence identified as SEQ ID NO:11, SEQ ID NO:12, or SEQ ID NO:13; analogues thereof wherein the amino acids in the sequence may be substituted, deleted or added, so long as the bone stimulatory activity in mammals derived the three dimensional structure of the sequence is

preserved; and conjugates of each of the polypeptides or analogues thereof, wherein if the polypeptide sequence contains a cysteine residue, there are at least two cysteine residues. Such a polypeptide can be substantially pure and the amino acid sequence can have a molecular weight in the range of from about 1000 to about 4000, or from about 1500 to about 3000, or from about 1500 to about 1800. The invention is also a first polypeptide that includes a sequence of amino acids sufficiently duplicative of a second polypeptide having an amino acid sequence corresponding such a polypeptide, or a functionally equivalent homologue thereof, such that the first polypeptide is encoded by a DNA that hybridizes under stringent conditions with DNA encoding the second polypeptide; or a chimeric bone stimulating factor including such an amino acid sequence; or an agent for use in prevention and treatment of a bone reduction related disease which includes such a polypeptide as an active ingredient; or a pharmaceutical composition for promoting bone growth, having a therapeutically effective amount of such a polypeptide; or a method of increasing bone growth in a mammal by administering a therapeutically effective amount of such a polypeptide; having an amino acid sequence of a polypeptide defined in claim 56 or 57; or the use of such a polypeptide for the treatment of osteoporosis or to promote bone growth in a mammal; or the use of such a polypeptide in the preparation of a medicament for use in promoting bone growth or the treatment of osteoporosis; or a diagnostic kit for determining the presence of such a polypeptide, the kit including an antibody to a the polypeptide linked to a reporter system wherein the reporter system produces a detectable response when a predetermined amount of the polypeptide and the antibody are bound together; or an antibody which binds to such a polypeptide, synthesized using the polypeptide; or an isolated DNA fragment which encodes the expression of any such polypeptide, or which differs from the fragment due to the degeneracy of the genetic code; or a vector including a DNA sequence which encodes the expression of any such polypeptide; or a process for producing such a polypeptide, which process includes:

- [0076] a) preparing a DNA fragment containing a nucleotide sequence which encodes the polypeptide;
- [0077] b) incorporating the DNA fragment into an expression vector to obtain a recombinant DNA fragment which contains the DNA fragment and is capable of undergoing replication;
- [0078] c) transforming a host cell with the recombinant DNA fragment to isolate a transformant which can express the polypeptide; and
- [0079] d) culturing the transformant to allow the transformant to produce the polypeptide and recovering the polypeptide from resulting cultured mixture.
- [0080] Finally, the invention includes an isolated DNA sequence encoding the amino acid sequence of any of the polypeptides of the invention, or an analogue thereof, wherein the amino acids in the sequence may be substituted, deleted or added, so long as bone stimulatory activity in mammals derived from the three dimensional conformation of the sequence is preserved in a polypeptide comprising the amino acid sequence; sequences which hybridize to the DNA and encode an amino acid sequence of a polypeptide which displays bone stimulatory activity in mammals; and DNA which differs from the sequence due to the degeneracy of the genetic code.

#### DESCRIPTION OF THE DRAWINGS

- [0081] Figure 1 shows the bone apposition rate ( $\mu\text{m}$  per day) for rats injected with 25 nmol (N=5 in both cases) of chemically synthesized polypeptides having the sequences of NAP-2 (SEQ ID NO:1) and NAP-2V (SEQ ID NO:2), respectively, compared to that of a group of control (N=5) rats. The error bars are  $\pm 1$  S.D.
- [0082] Figure 2 graphically illustrates the observed bone mineral apposition rate ( $\mu\text{m}$  per day) for rats injected with chemically synthesized polypeptides of A, NAP-2V-(1-26) (SEQ ID NO:11); B, NAP-2V-(13-26; gln 25 $\rightarrow$ glu25) (SEQ ID NO:12, ), C, NAP-2V-(10-26) (SEQ ID NO:13), D, NAP-2V-(11-26) (SEQ ID NO:14), and E, NAP-2V-(12-26) (SEQ ID NO:15). The first bar in the graph

represents the control. The number of rats used for the determinations were 4, 4, 3, 4, 4, and 4, respectively. The error bars are  $\pm 1$  S.D.

[0083] Figure 3 graphically illustrates the observed bone mineral apposition rate ( $\mu\text{m}$  per day) for rats injected with chemically synthesized polypeptides having SEQ ID NO:17 (second bar), SEQ ID NO:18 (third bar) compared to a control (first bar). The error bars are  $\pm 1$  S.E.

[0084] Figure 4 graphically illustrate the dose dependency of the bone mineral apposition rate ( $\mu\text{m}$  per day) found for rats injected with the indicated amounts of polypeptide having SEQ ID NO:18. The error bars are  $\pm 1$  S.E.

[0085] Figure 5 illustrates the amino acid sequences of polypeptides tested, corresponding amino acids aligned with each other, the active peptides being shown above the line and sequences which were not found to stimulate bone growth being below the line. Approximate molecular weights are shown below the sequence identification numbers.

#### MATERIALS, METHODS AND RESULTS

[0086] Polypeptides having the sequences of NAP-2 (SEQ ID NO:1) and NAP-2V (SEQ ID NO:2) were chemically synthesized directly according to standard methods and experiments were conducted to determine whether the chemically synthesized polypeptides displayed activity.

[0087] Experiments were conducted simultaneously on three groups of male Sprague-Dawley rats, there being five rats in each group. Each rat weighed between 250 and 350 g. Each rat of the first group was given, by subcutaneous injection into the left gluteus maximus region, 200  $\mu\text{l}$  of a 1% aqueous acetic acid solution containing 25 nmol (about 191  $\mu\text{g}$ ) of NAP-2 (SEQ ID NO:1). Each rat of the second group was similarly given 200  $\mu\text{l}$  of a 1% aqueous acetic acid solution containing 25 nmol (about 207  $\mu\text{g}$ ) of NAP-2V (SEQ ID NO:2). Each rat of the third group, the control group, was similarly given 200  $\mu\text{l}$  of 1% acetic acid solution.

[0088] Immediately following administration of the test solution, 300  $\mu\text{l}$  of an aqueous solution of tetracycline hydrochloride was administered intramuscularly into the right gluteus maximus, the concentration of

tetracycline being sufficient to obtain a dosage of about 24mg/kg of rat body weight. A second dose of tetracycline hydrochloride solution was administered about 48 hours after the first dose. The rats were sacrificed about 24 hours after administration of the second dose of tetracycline.

[0089] Sections of the lower metaphysis of the right femur were used for bone measuring the bone mineral apposition rate. Processing of the bone material for measurement has been described previously. See, for example, international patent application No. PCT/CA 94/00144 published under No. WO 94/20615 on September 15, 1994. The results obtained are summarized in Table One and Figure 1.

TABLE ONE: Comparison of the Group Arithmetic Means of Bone Apposition Rates ( $\mu\text{m}/\text{day}$ ) Among Groups Administered with NAP-2, NAP-2V and control solutions shown in Figure 1			
	Control	SEQ ID NO:1	SEQ ID NO:2
Mean	0.99 $\mu\text{m}/\text{d}$	1.23 $\mu\text{m}/\text{d}$	1.28 $\mu\text{m}/\text{d}$
S.D.	0.04	0.05	0.08
N	5	5	5
	t	d.f.	p
Control Group vs SEQ ID NO:1	7.91	8	<0.001
Control Group vs SEQ ID NO:2	7.03	8	<0.001
SEQ ID NO:1 vs SEQ ID NO:2	1.15	8	>0.20

[0090] Polypeptides having the sequence corresponding to either SEQ ID NO:1 or SEQ ID NO:2 have thus been found to stimulate bone growth.

[0091] In a second set of experiments, twenty-four male Sprague-Dawley rats (Charles River Laboratory) were divided into six groups of four. Each of the

first group, the control group, was injected in the right thigh with 200  $\mu$ l of 0.1% acetic acid solution. The rats of the other groups were each injected in the right thigh with about 200  $\mu$ l of a 0.1% acetic acid solution containing about 25 nmol of a chemically synthesized polypeptide as follows:

Group	Polypeptide	SEQ ID NO
A	NAP-2V-(1-26)	11
B	NAP-2V-(13-26; gln <sup>25</sup> →glu <sup>25</sup> )	12
C	NAP-2V-(10-26)	13
D	NAP-2V-(11-26)	14
E	NAP-2V-(12-26)	15

[0092] Immediately after injection of the polypeptide (control) solution, each rat was injected in the right gluteus maximus with 200  $\mu$ l of a 1M tetracycline hydrochloride (Sigma Chemical) solution. This dosage of tetracycline is about 16 mg per kg of animal body weight.

[0093] About 48 hours later, each rat was injected in the left gluteus maximus with the same dosage of tetracycline. Twenty-four hours later, the rats were sacrificed by CO<sub>2</sub> narcosis and the right femur taken for bone mineral apposition rate determination. One rat died during the course of the experiments.

[0094] Immediately after dissection, a bone sample was fixed in 10% formaldehyde solution at pH 7.4. Later the same day, a 1:1 H<sub>2</sub>O-acetone solution was exchanged for the formaldehyde solution. This was exchanged twice the following day with acetone. This was exchanged the following day by a 1:1 acetone-Spurr's medium solution, which was exchanged later the same day with Spurr's medium. The following day each sample was embedded in a fresh change of Spurr's medium and cured at 60°C for 24 hours, followed by curing at 80°C for 24 hours.

[0095] Each cured block was cut into 400  $\mu$ m thick sections using a Leitz saw microtome equipped with a diamond charged blade. The relatively thick sections were ground down between two ground glass plates pre-roughened



with carborundum powder to a final thickness of about 10  $\mu\text{m}$ , water being used as the grinding lubricant. The thin sections were dried and mounted unstained in Permount (Fisher).

[0096] Measurements were made using a Leitz scanning light microscope photometer MPV-CD magnifying the sections 16X, as described in international patent application No. PCT/CA94/00144.

[0097] The results obtained are shown in Table Two and Figure 2.

<b>Table TWO: Comparison of Group Arithmetic Means of Bone Apposition Rates (<math>\mu\text{m}</math> per day) shown in Figure 2.</b>						
	Control	Group A	Group B	Group C	Group D	Group E
Mean	1.08	1.47	1.70	1.18	1.07	1.04
S.D.	0.08	0.05	0.09	0.08	0.07	0.08
N	4	4	3	4	4	4

	t	d.f.	p
Control vs Group A	7.90	6	<0.001
Control vs Group B	9.69	5	<0.001
Group A vs Group B	4.87	5	

[0098] The polypeptide having the sequence identified as SEQ ID NO:12 has two sites susceptible to cleavage by the peptidase plasmin, between the lysine-threonine and the lysine-asparagine residues, respectively. In a third set of experiments, nine female Sprague-Dawley rats (average weight, about 300 gm) were divided into three groups of three. Each of the first group, the control group, was injected with 400  $\mu\text{l}$  of buffer solution, 50 mM sodium acetate adjusted to pH 4.5 followed immediately by 200  $\mu\text{l}$  of a 1M tetracycline hydrochloride solution. The rats of the second group were each injected with 400  $\mu\text{l}$  of buffer containing 100 nmoles of a chemically synthesized polypeptide having the sequence identified as SEQ ID NO:17 followed immediately by 200  $\mu\text{l}$  of a 1M tetracycline hydrochloride solution. The rats of

the third group were each injected with 400  $\mu$ l of buffer containing 100 nmoles of a chemically synthesized polypeptide having the sequence identified as SEQ ID NO:18 followed immediately by 200  $\mu$ l of a 1M tetracycline hydrochloride solution.

[0099] After about 48 hours, a second dose of tetracycline hydrochloride was administered to each rat. After about another 24 hours, rats were sacrificed by carbon dioxide narcosis.

[00100] The right femur of each animal was taken and treated, as described above, and measurements made as described above. The results obtained are shown in Table Three and Figure 3.

TABLE THREE: Comparison of the Group Arithmetic Means of Bone Apposition Rates ( $\mu$ m/day) Among Groups Administered with polypeptides having SEQ ID NO:17, SEQ ID NO:18, control solutions shown in Figure 3			
	Control	SEQ ID NO:17	SEQ ID NO:18
Mean	1.256 $\mu$ m/d	1.859 $\mu$ m/d	2.406 $\mu$ m/d
S.E.	0.06	0.10	0.25
N	3	3	3

[00101] In a fourth set of experiments, dose dependence of the measured bone mineral apposition rate on the amount of polypeptide having SEQ ID NO:18 was evaluated. Twenty-eight female Sprague-Dawley rats (average weight, about 300 gm) were divided into seven groups of four. The rats were treated as described in the previous set of experiments except that the amount of test polypeptide varied as follows: 25, 50, 100, 200, 400 and 800 nmoles. The results obtained are shown in Table Four and Figure 4.

Table FOUR: Dose Dependency of Bone Apposition Rates ( $\mu\text{m}/\text{day}$ ) on Amount of Polypeptide having SEQ ID NO:18							
	Control	25 nmol	50 nmol	100 nmol	200 nmol	400 nmol	800 nmol
Mean	1.320	1.686	1.753	1.907	2.209	2.123	2.139
S.E.	0.04	0.08	0.03	0.03	0.08	0.06	0.03
N	4	4	4	4	4	4	4

[00102] As graphically illustrated in Figure 2, NAP-2V-(1-26) (SEQ ID NO:11) and NAP-2V-(13-26; gln<sup>25</sup>→glu<sup>25</sup>) (SEQ ID NO:12) act to stimulate bone growth, the latter of these two polypeptides displaying greater activity. As illustrated in Figure 3, NAP-2V-(15-21) having its amino and carboxy termini protected (SEQ ID NO:18) has greater bone stimulatory activity than the similarly protected NAP-2V-(13-26; gln<sup>25</sup>→glu<sup>25</sup>) (SEQ ID NO:17).

[00103] NAP-2V-(10-26) (SEQ ID NO:13) appeared to cause a small increase in the observed bone apposition rate, although the significance of the observed increase was questionable. NAP-2V-(11-26) (SEQ ID NO:14) and NAP-2V-(12-26) (SEQ ID NO:15) were found to have no effect on observed bone mineral apposition rate. The sequence of NAP-2V-(10-26) retains both the cys<sup>10</sup> and cys<sup>12</sup> residues. The sequences of NAP-2V-(11-26) and NAP-2V-(12-26) each retain the cys<sup>12</sup> residue. The sequence of NAP-2V-(13-26; gln<sup>25</sup>→glu<sup>25</sup>) retains neither of the cys<sup>10</sup> and cys<sup>12</sup> residues. All of these NAP-2V subsequences lack the cys<sup>36</sup> and cys<sup>52</sup> residues present in the parent NAP-2V. It may be that the reduced activity of NAP-2V-(10-26), NAP-2V-(11-26) and NAP-2V-(12-26) is due to spontaneous intermolecular disulfide bonding that prevents a polypeptide-receptor interaction required for the bone stimulatory effect, but this is not known for certain.

[00104] The sequence of NAP-2V-(13-26; gln<sup>25</sup>→glu<sup>25</sup>) is different from the corresponding subsequence of NAP-2V at the 25 position, a glutamic acid residue being present in place of the glutamine residue. It would of course be

expected that the subsequence having the glutamine residue as occurs in NAP-2V would also act to stimulate bone growth in mammals.

[00105] It has been postulated that NAP-2 contains two internal disulfide bonds, between Cys-5 and Cys-31, and Cys-7 and Cys-47, respectively (Baggiolini, M., Clemetson, K.J., Walz, A. International Patent Application No. PCT/EP89/01389, published June 14, 1990 under WO90/06321.). By extension, sequences and subsequences disclosed herein that contain the corresponding cysteine residues would likely contain similar linkages therebetween.

[00106] It will of course be understood, without the intention of being limited thereby, that a variety of other substitutions of amino acids is possible while preserving the structure responsible for the bone stimulatory effect of the subsequences of NAP-2V disclosed herein. Conservative substitutions are described in the patent literature, as for example, in United States Patent No. 5,264,558. It is thus expected, for example, that interchange among non-polar aliphatic neutral amino acids, glycine, alanine, proline, valine and isoleucine, would be possible. Likewise, substitutions among the polar aliphatic neutral amino acids, serine, threonine, methionine, asparagine and glutamine could possibly be made. Substitutions among the charged acidic amino acids, aspartic acid and glutamic acid, could probably be made, as could substitutions among the charged basic amino acids, lysine and arginine. Substitutions among the aromatic amino acids, including phenylalanine, histidine, tryptophan and tyrosine would also likely be possible. These sorts of substitutions and interchanges are well known to those skilled in the art. Other substitutions might well be possible. A peptide containing an amino acid sequence that can be aligned with that of SEQ ID NO:11, SEQ ID NO:12 or NO:18 and having less than 100% homology therewith may retain at least part of the bone stimulating effect thereof. Of course, it would also be expected that the greater percentage of homology, say 70%, 80%, 90%, or more, could increase the degree of retained bone stimulating activity.

[00107] "Sequence identity or homology", as used herein, refers to the sequence similarity between two polypeptide molecules or between two

nucleic acid molecules. When a position in both of the two compared polypeptide sequences, for example, is occupied by the same amino acid (for example, if a position in each of two polypeptide molecules is an alanine residue, then the molecules are homologous or sequences are identical at that position. The percent of homology between two molecules or sequence identity between two sequences is a function of the number of such matching positions shared by the two sequences divided by the number of positions compared x 100. For example, if 6 of 10, of the positions in two sequences are the same then the two sequences are 60% homologous or have 60% sequence identity. By way of example, the polypeptide sequences METLIA and MPTWIF share 50% homology or sequence identity. Generally, a comparison is made when two sequences are aligned to give maximum homology.

[00108] The comparison of sequences and determination of percent homology between two sequences can be accomplished using a mathematical algorithm. In this specification, the alignment was performed according to two methods, the Clustal method and the J. Hein method. Of these, the Clustal method is preferred.

[00109] The Clustal algorithm (as applied here using software available from DNASTAR Inc., 1228 South Park Street, Madison, Wisconsin, USA, 1994) is recommended for aligning sequences whose similarity might not necessarily be evolutionary. The algorithm is described by Higgins, D.G. *et al.* 1989. *CABIOS* 5:151. The same software programme provides for aligning sequences according to the Jotun Hein method, which is recommended for aligning sequences of highly evolved families that have clear evolutionary relationship. The algorithm is described by Hein, J. 1990. *Methods in Enzymology* 183:626. Programme default settings (standard parameters) were used. In the case of weighting amino acid residues based on evolutionary substitution patterns, charge, structural and chemical similarity, the default PAM250 setting was used. For protein alignments, the pairwise alignment parameters are Ktuple =1, Gap penalty =3, Window =5, and Diagonals Saved =5 were used.

[00110] Insofar as deletion of one or more amino acids is concerned, it is likely that deletions of a small number of amino acids from each end of either sequence might be possible, bearing in mind the observation that the deletions to obtain SEQ ID NOs: 14 and 15 yield polypeptides which do not appear to enhance bone growth. Further, symmetrical, or nearly symmetrical deletions would likely be the most possible to be made while retaining the three-dimensional configuration. Internal deletions, although likely to be possible to some limited extent, should be few.

[00111] Additions of amino acids could very likely be made at the ends of the sequence, and as with deletions, symmetrical or nearly symmetrical additions to the carboxy and amino terminals are likely to be possible. Internal additions, although likely to be possible to some limited extent, should be few.

[00112] Of the above-listed modifications to the sequence, terminal additions are most likely to be most useful, as such a modification can serve a variety of functions: an identifying group as for use in a radioimmunoassay; or a linking group, as examples.

[00113] A polypeptide of the present invention can be improved with respect to possible degradation, as might occur in the body in the presence of protease, for instance, by protection of the C-terminus, the N-terminus, or both the C-terminus and N-terminus of the polypeptide.

[00114] As used herein, "protected" terminal amino group refers to a terminal amino group (N-terminus) coupled with any of various amino-terminal protecting groups that can be employed in peptide synthesis. Examples of suitable groups include acyl protecting groups, for example, formyl, acetyl, benzoyl, trifluoroacetyl, succinyl, and methoxysuccinyl; aromatic urethane protecting groups, for example benzyloxycarbonyl; and aliphatic urethane protecting groups, for example *t*-butoxycarbonyl or adamantyloxycarbonyl (Gross and Mienhofer, eds., *The Peptides*, vol 3, pp. 3 to 88 (Academic Press, New York, 1981)).

[00115] As used herein, "protected" terminal carboxyl group refers to a terminal carboxyl group (C-terminus) coupled with any of various

carboxy-terminal protecting groups. As will be readily apparent to a person skilled in the art, suitable groups include *t*-butyl, benzyl or other acceptable groups linked to the terminal carboxyl group through an ester or ether bond.

[00116] Compounds within the scope of this invention can be synthesized chemically by means well known in the art such, for example, solid phase peptide synthesis. The synthesis is commenced from the carboxy-terminal end of the peptide using an  $\alpha$ -amino protected amino acid. *t*-Butyloxycarbonyl (Boc) protective groups, or other suitable protective groups, can be used (Stewart *et al.*, "Solid-Phase Peptide Synthesis," W. H. Freeman Co., San Francisco (1969); Merrifield, *J. Am. Chem. Soc.* 85:2149-2154 (1963); Vale *et al.*, *Science* 213, 1394-1397 (1981), and Marke *et al. J. Am Chem. Sci.* 103, 3178 (1981)). Synthetic methods are also described in "Principles of Peptide Synthesis" M. Bodansky Ed. (Spring-Verlag 1984). These and other methods of peptide synthesis are also exemplified by U.S. Patent Nos. 3,862,925, 3,842,067, 3,972,859, 4,105,602, 4,683,291, 4,244,946 and 4,305,872.

[00117] Compounds may also be synthesized using manual or automatic techniques, for example, an Applied BioSystems 430A Peptide Synthesizer (Foster City, California) or a Biosearch SAM 11 automatic peptide synthesizer (Biosearch, Inc., San Rafael, California).

[00118] A compound "derived from" a polypeptide having a particular amino acid sequence is any molecular entity which is identical, substantially homologous, or otherwise functionally or structurally equivalent to that polypeptide. Thus, a molecule derived from a particular polypeptide may encompass the amino acid sequence of the polypeptide, any portion of that polypeptide, or other molecular entity that functions to stimulate bone growth. A molecule derived from such a binding domain will mimic the polypeptide from which it is derived. Such molecular entities may include peptide mimetics and the like.

[00119] "Peptides mimetics" are structures which serve as substitutes for peptides in interactions with acceptor molecules (see Morgan *et al.* (1989) *Ann. Reports Med. Chem.* 24:243-252 for a review of peptide mimetics).

Peptide mimetics, as used herein, include synthetic structures which may or may not contain amino acids and/or peptide bonds, but retain structural and functional features of a peptide from which they are derived. The term, "peptide mimetics" also includes peptoid and oligopeptoids, which are peptides or oligomers of N-substituted amino acids (Simon et al. (1972) *Proc. Natl. Acad. Sci USA* 89:9367-9371). Further included as peptide mimetics are peptide libraries, which are collections of peptides designed to be a given amino acid length and representing all conceivable sequences of amino acids corresponding thereto.

[00120] Two polypeptide sequences are "substantially homologous" when at least about 85% (preferably at least about 85% to 90%, and most preferably at least about 95%) of the nucleotides or amino acids match over a defined length of the polypeptide. As used herein, substantially homologous also refers to sequences showing identity to the specified polypeptide sequence.

[00121] Peptide mimetics which structurally and functionally mimic the polypeptides having bone stimulatory activity described herein, will also find use herein and may be generated using the following strategies and procedures. Generally, mimetics are designed based on information obtained by systematic replacement of L-amino acids by D-amino acids, replacement of side chain moieties by a methyl group or pseudoisosteric groups with different electronic properties (see Hruby et al. (1990) *Biochem, J.* 268:249-262), and by systematic replacement of peptide bonds in the above described peptide inhibitors with amide bond replacements. For example, analogues containing amide bond surrogates may be used to investigate aspects of peptide structure and function, such as rotational freedom in the backbone, intra and intermolecular hydrogen-bond patterns, modifications of local and total polarity and hydrophobicity, and oral bioavailability.

[00122] Local conformational constraints can also be introduced to determine conformational requirements for activity of a potential peptide mimetic having bone stimulatory activity. For example,  $\beta,\beta$ -distributed amino acids may be used to examine the effects of conformational constraints on



peptide activity (see, e.g. Manning et al. (1982) *J. Med. Chem.* 25:408-414; Mosberg et al. (1983) *Proc. Natl. Acad. Sci. USA* 106:506-512; Pelton et al. (1985) *Proc. Natl. Acad. Sci. USA* 82:236-239).

[00123] The mimetics can include isosteric amide bonds such as  $\psi$ [CH<sub>2</sub>S],  $\psi$ [CH<sub>2</sub>NH],  $\psi$ [CNH<sub>2</sub>],  $\psi$ [NHCO],  $\psi$ [COCH<sub>2</sub>] and  $\psi$ [(E) or (Z) CH=CH] see, for review, Spatola (1983) in "Chemistry and Biochemistry of Amino Acids, Peptides and Proteins," Volume VII. (Weinstein ed.), Marcel Dekker, New York, 267-357). The synthetic molecules can also include D-amino acids to stabilize or promote reverse turn conformations and to help stabilize the molecule from enzymatic degradation (see, e.g. Freidinger et al (1985) in "Peptides: Structure and Function." (Deber et al. eds.), Pierce Chem Co., Rockford, Ill., 549-552; Sawyer et al (1980) *Proc. Natl. Acad. Sci. USA* 77:5754-5758; Torchiana et al (1978) *Arch. Int. Pharmacol. Ther.* 235:170-176). Cyclic amino acid analogues may be used to constrain amino acid residues to particular conformational states, e.g.  $\alpha\alpha'$ - and  $\beta\beta$ - substituted cyclic amino acids such as 1-aminocyclopentanecarboxylic acid (cycloleucine) and  $\beta,\beta$ - cyclopentamethylene- $\beta$ - mercaptopropionic acid (see Hruby et al (1990), *supra*).

[00124] The mimetics can also include mimics of polypeptide secondary structure—structures which can model the 3-dimensional orientation of amino acid residues into the known secondary conformations of proteins—including  $\beta$ - turn mimetics, such as phenoxathin ring system, and  $\beta$ - sheet mimics, such as epindolidione structures. Design synthesis and conformational analysis of a  $\alpha$ -helix inducing template has been described (Kemp et al (1988) *Tetrahedron Lett.* 29:4931; Kemp et al. (1988) *Tetrahedron Lett.* 29:4935).

[00125] A potential mimetic can be tested, or pre-screened, for potential activity as a bone stimulating compound by measuring the affinity of the compound for an antibody raised against the polypeptide from which the mimetic is derived. As described above for polypeptides, those mimetics that react positively with the antibody to the already known peptide could then be tested for bone stimulatory effects *in vivo* using the system described her in

for rats, for example. Antibodies raised against a polypeptide having the amino acid sequence identified as SEQ ID NO:9 would be particularly useful in this context.

[00126] Peptoids will find use herein. Peptoids are oligomers of N-substituted amino acids (Simon et al (1972), *supra*). and can be used as motifs for the generation of chemically diverse libraries of novel molecules, which can then be tested for binding and bone stimulatory activity. The monomers may incorporate t-butyl-based side-chain and 9-fluorenylmethoxy-carbonyl  $\alpha$ -amine protection. Oligomerization of the peptoid monomers may be performed by for example, *in situ* activation by either benzotriazol-1-yloxytris(pyrrolidino)phosphonium hexafluorophosphate or bromotris(pyrrolidino)phosphonium hexafluorophosphate. Other steps are identical to conventional peptide synthesis using  $\alpha$ -(9-fluorenylmethoxycarbonyl)amino acids. Oligopeptoids may be identified which have affinities comparable to the corresponding polypeptides and, thus, are potentially useful as bone stimulatory agents.

[00127] Compounds of the present invention and compositions containing them find use in numerous therapeutic and prophylactic applications in the prevention and treatment of bone reduction related to a disease. Compounds can thus be used as treatments to promote bone growth, in the treatment of osteoporosis, for example, by any suitable route. The preferred routes are suitable for delivery of polypeptide-type compounds to the bloodstream of a subject, bearing in mind proper storage and handling conditions required for polypeptides such as those described herein.

[00128] Thus the present invention also provides compositions containing an effective amount of compounds of the present invention, including the nontoxic addition salts, amides and esters thereof, which may, alone, serve to provide the treatment benefits described above. Such compositions can also be provided together with physiologically tolerable liquid, gel or solid diluents, adjuvants and excipients.

[00129] In the above examples involving subsequent examples of NAP-2V, about 75 nmol of polypeptide per kg of bodyweight of animal was used per

administration. In practice, particularly as human subjects are concerned, the daily dosage may well be between 0.01 and 300 mg or more per kg of bodyweight. More preferably, the dosage would be in the neighborhood of from about 0.1 to about 30 mg per kg of bodyweight. It may be that the preferred frequency of administration would be greater or less than once per day, depending upon the route of administration, convenience, and the variation of effectiveness of treatment with frequency of and amount used per administration. The dosage administered also depends on the subject and to which effect such administration is to give. The dosage of any one or more of the compounds will depend on many factors including the specific compound or combination of compounds being utilized, the mode of administration, and the mammal being treated. Dosages of a particular compound or combination of compounds can be determined using conventional considerations; for example, by customary comparison of the differential activities of the subject compounds and that of a known agent, that is, by means of an appropriate pharmacological protocol in which, for example, bone density of subjects is measured over time.

[00130] Pharmaceutical preparations include any of the compounds prepared as an injectable solution, including an injectable solution prepared just prior to use, for promoting bone growth and/or treatment of osteoporosis. An injectable can be either a liquid solution or suspension; solid forms suitable for solution in, or suspension in, liquid prior to injection may also be prepared. The preparation may also be emulsified. The active polypeptide is often mixed with diluents and excipients which are physiologically tolerable and compatible with the polypeptide. Suitable diluents and excipients are, for example, water, saline, dextrose, glycerol, or the like, and combinations thereof. In addition, if desired, the compositions can contain minor amounts of auxiliary substances such as wetting or emulsifying agents, stabilizing or pH buffering agents, and the like.

[00131] Pharmaceutical preparations include the employment of the compounds in admixture with conventional excipients, that is, pharmaceutically acceptable organic or inorganic carrier substances which do

not deleteriously react with the compounds, and which possibly enhance the storage and handling stability of the compounds. The preparative procedure may include the sterilization of the pharmaceutical preparations. The compounds may be mixed with auxiliary agents such as lubricants, preservatives, stabilizers, salts for influencing osmotic pressure, etc., which do not react deleteriously with the compounds.

[00132] The compositions are conventionally administered parenterally, by injection, for example either subcutaneously or intravenously. Additional formulations which are suitable for other modes of administration include suppositories, intranasal aerosols, and, in some cases, oral formulations. For suppositories, traditional binders and excipients may include, for example, polyalkylene glycols or triglycerides; such suppositories may be formed from mixtures containing the active ingredient in the range of 0.5% to 10%, preferably 1%-2%. Oral formulations include such normally employed excipients as, for example, pharmaceutical grades of mannitol, lactose, starch, magnesium stearate, sodium saccharin, cellulose, magnesium carbonate, and the like. These compositions take the form of solutions, suspensions, tablets, pills capsules, sustained release formulations, or powders, and contain 10% - 95% of active ingredient, preferably 25% - 70%. These oral formulations include formulations designed to protect the peptide until it can be absorbed.

[00133] The peptide compounds may be formulated into the compositions as neutral or salt forms. Pharmaceutically acceptable non-toxic salts include the acid addition salts (formed with the free amino groups) and which are formed with inorganic acids such as, for example, hydrochloric or phosphoric acids, or such organic acids as acetic, oxalic, tartaric, mandelic, and the like. Salts formed with the free carboxyl groups may be derived from inorganic bases such as, for example, sodium, potassium, ammonium, calcium, or ferric hydroxides, and such organic bases as isopropylamine, trimethylamine, 2-ethylamino ethanol, histidine, procaine, and the like.

[00134] The compounds of the invention can be homopolymerized to themselves (i.e., (peptide)<sub>n</sub>) or, heteropolymerized to one another. The

compounds can also be conjugated to biocompatible polymeric compounds, such as BIOPOL™ (WR Grace & Co.-Conn.).

[00135] If prepared using recombinant techniques, a DNA sequence encoding a desired polypeptide of the present invention is synthesized using standard automated techniques, or the coding sequences or portions thereof are retrieved from cDNA or genomic libraries. This DNA is ligated into suitable expression vectors and these vectors are transformed into appropriate hosts. A variety of expression vector/host cell systems can be used, including both procaryotic and eukaryotic culture systems.

[00136] Procaryotes most frequently are represented by various strains of *E. coli*. However, other microbial strains may also be used, such as bacilli, for example *bacillus subtilis*, various species of *Pseudomonas*, or other bacterial strains. In such procaryotic systems, plasmid vectors which contain replication origins, and control sequences derived from a species compatible with the host are used. For example, *E. coli* is typically transformed using derivatives of pBR322, a plasmid derived from an *E. coli* species (Bolivar et al., (1977) *Gene* 2:95. Commonly used procaryotic control sequences, which are defined herein to include promoters for transcription initiation, optionally with an operator, along with ribosome binding site sequences, include such commonly used promoters as the beta-lactamase (penicillinase), lactose (lac) promoter systems (Chang et al., (1977) *Nature* 198:1056), the tryptophan (trp) promoters system (Goeddel et al., (1990) *Nucleic Acids Res* 8:4057), and the lambda-derived P<sub>L</sub> promoter and N-gene ribosome binding site (Shimatake et al., (1981) *Nature* 292:128). However, any available promoter system compatible with procaryotes can be used.

[00137] The expression systems useful in the eukaryotic systems of the invention comprise promoters derived from appropriate eukaryotic genes. A class of promoters useful in yeast, for example, include promoters for synthesis of glycolytic enzymes, including alcohol dehydrogenase promoters, glyceraldehyde-3-phosphate dehydrogenase promoter (Holland & Holland, (1980) *J Biol Chem* 25:2596), alpha-factor promoter (Bitter et al., (1984) *Proc Natl Acad Sci* 81:5330), the gal promoter (Johnston & David, (1984) *Mol*

*Cell Biol* 4:1440) those for 3-phosphoglycerate kinase (Hitzeman et al., (1980) *J. Biol Chem* 256:1385) or the Leu2 gene obtained from YEp13 (Broach, J., et al., (1978) *Gene* 8:121).

[00138] Suitable mammalian promoters include the early and late promoters from SV40 (Fiers et al., (1978) *Nature* 273:113) or other viral promoters such as those derived from polyoma, adenovirus II, bovine papilloma virus or avian sarcoma viruses. Suitable viral and mammalian enhancers are cited above. In the event plant cells are used as an expression system, the nopaline synthesis promoter is appropriate (Depicker, A., et al., (1982) *J Mol Appl Gen* 1:56).

[00139] The expression systems are included on replication vectors or are caused to integrate into the chromosome of a recombinant host. For systems wherein the vectors include a replication system, these may be low or high copy number, usually having copy numbers of fewer than about 1000, although in certain situations, runaway vectors may be employed. Whether provided on a vector intended for integration or in a replication system, the sequence encoding a polypeptide of the invention may be ligated in tandem with an amplifying gene such as dihydrofolate reductase, metallothioneins, thymidine kinase, or the like. In procaryotic systems, both the amplifying gene and the target gene can be under the regulation of the same transcriptional and translational regulatory regions.

[00140] Usually, the vector will include a marker which allows for selection of host cells containing the expression system; the nature of these markers depends on the host and is understood in the art. In addition to required regulators such as promoters, additional sequences such as enhancers can also be employed to enhance the level of transcription. If the polypeptide is to be secreted, an upstream sequence encoding signal peptides such as those described in U.S. Pat. Nos. 4,336,336; 4,338,397; and 4,546,082 may be employed. The signal sequence is enzymatically cleaved as the polypeptide product is secreted.

[00141] Depending on the host cell used, transformation is done using standard techniques appropriate to such cells. The calcium treatment

employing calcium chloride, as described by Cohen, S.N., (1972) *Proc Natl Acad Sci USA* 69:2110; or the RbCl method described in Maniatis et al., *Molecular Cloning: A Laboratory Manual* (1982) Cold Spring Harbor Press. p. 254 is used for procaryotes or other cells which contain substantial cell wall barriers. Infection with *Agrobacterium tumefaciens* (Shaw, C.H., (1938) et al., *Gene* 23:315) is used for certain plant cells. For mammalian cells without such cell walls, the calcium phosphate precipitation method of Graham and van der Eb, (1978) *Virology* 52:2:546 is preferred. Transformations into yeast are carried out, for example, according to the method of Van Solingen, P., et al., (1977) *J Bacter* 130:946; and Hsiao, C.L., et al., (1979) *Proc Natl Acad Sci USA* 76:3829.

[00142] In general, after construction of a suitable expression system, the system is transfected into the appropriate host and successful transformants are selected by markers contained on the expression vectors. Successfully transformed colonies are then cultured in order to produce the desired polypeptide. It is sometimes preferred that a promoter which can be controlled by regulating conditions in the environment be used so that the cells can be grown under conditions where the gene encoding the desired polypeptide of the invention is not expressed, and then production of the polypeptide induced by appropriate manipulation of conditions. For example, if the *trp* promoter is used in *E. coli*, the cells are grown in the presence of tryptophan and expression is then induced by diminution of tryptophan concentration or by addition of a tryptophan analogue such as indolylacetic acid. If the gene is under control of the PL promoter, the cells are grown at relatively low temperature, such as at about 35°C., to a suitable cell density, and the temperature is then elevated to activate this promoter. If produced in bacterial hosts as a mature intracellular polypeptide, the N-terminal methionine may or may not be cleaved. In mammalian systems, for example, the use of the metallothionein promoter permits induction by addition of heavy metals or glucocorticoids. This protocol is preferred to prevent premature accumulation of the polypeptide which might be harmful to the growth of the cell.

[00143] The polypeptide can be produced intracellularly, or in secreted form by construction of vectors in which the peptide is preceded by a signal peptide workable in the appropriate host.

[00144] The polypeptide is recovered from the medium or from the cells using suitable techniques generally known in the art, and purified by, for example, ion exchange chromatography, ammonium sulfate precipitation, gel permeation chromatography, and so forth.

[00145] The polypeptide made available by the invention disclosed herein can be used to obtain antisera thereto (Stites, D.P. and A.I. Terr. 1991. In Basic & Clinical Immunology, 7th Ed. Appleton and Lange, Norwalk, Connecticut and San Mateo California). Methodology and products can be developed using an antibody to a polypeptide for use in detecting the polypeptide with which the antibody binds. This apparently having been accomplished at least for the polypeptide having the sequence of CTAP-III (SEQ ID NO:3) (Baggiolini, M., Clemetson, K.J., Walz, A. International Patent Application No. PCT/EP89/01389, published June 14, 1990 under WO90/06321). Methodology and products can be developed using an antibody to a polypeptide for use in detecting the polypeptide with which the antibody binds.

[00146] For example, an antibody can be linked to or conjugated with a reporter system which is set up to indicate positively binding of the polypeptide to the antibody. Well known reporter systems include radioimmuno assays (RIAs) or immunoradiometric assays (IRMAs). Alternatively, an enzyme-linked immunosorbent assay (ELISA) would have in common with RIAs and IRMAs a relatively high degree of sensitivity, but would generally not rely upon the use of radioisotopes. A visually detectable substance may be produced or at least one detectable in a spectrophotometer. An assay relying upon fluorescence of a substance bound by the enzyme being assayed could be used. It will be appreciated that there are a number of reporter systems which may be used, according to the present invention, to detect the presence of a particular polypeptide. With



standardized sample collection and treatment, polypeptide presence above a threshold amount in blood serum could well be determined.

**[00147]** Such an antibody-linked reporter system could be used in a method for determining whether blood serum of a subject contains a deficient amount of the polypeptide. Given a normal threshold concentration of such a polypeptide in blood serum of a given type of subject, test kits could thus be developed.

**[00148]** A further advantage may be obtained through chimeric forms of the protein, as known in the art. A DNA sequence encoding the entire protein, or a portion of the protein, could thus be linked with a sequence coding for the C-terminal portion of *E. coli*  $\beta$ -galactosidase to produce a fusion protein, for example. An expression system for human respiratory syncytial virus glycoproteins F and G is described in United States Patent No. 5,288,630 issued February 22, 1994 and references cited therein, for example.

**[00149]** All references cited in this specification are incorporated herein by reference, including United States Provisional Patent Application Serial No. 004,314 filed September 26, 1995.